



ICGCW 2025

**7th INDIAN CANCER GENETICS CONFERENCE & WORKSHOP
Tata Memorial Hospital & ACTREC, Mumbai, India**



ICGC2025 CONFERENCE TMH, PAREL: 22-23 Feb'25 (Saturday & Sunday)

**THREE PRECONFERENCE
HANDS ON WORKSHOPS
20-21st Feb 2025**

- Genetic Counselling of Common Hereditary Cancers: TMH, Mumbai
- NGS Wetlab, DryLab & Reporting: ACTREC, Navi Mumbai
- Functional Genomics & Tissue Culture, ACTREC, Navi Mumbai

Deep Dive To know More about Less Known Genes

Navigating The Rising Tide of VUS

*My
Rare Gene Diary*



- Should Germline or Somatic report include these genes / variants ?
- How should they be described ?
- How to act (or not) on these ?

Registration Link
for a Deep Dive to learn together & reach consensus

ALL ABOUT LESS KNOWN or MODERATE RISK GENES, HYPOMORPHIC ALLELES & VUS

TMC Cancer Genetics Unit has CURATED germline genotype & phenotype data of 10,000+ cases - not only well characterised genes like BRCA1/2, MLH1, MSH2, TP53, PALB2) but also Less known genes (CHEK2, BRIP1, RAD51, CDH1, MSH6, PMS2, SDH, & many more We have reclassified hundreds of ClinVar VUS as Likely Benign & a few as Likely Pathogenic

Registration fee (with 18% GST) *Will be Half for Trainees & GCs & *Double for Industry delegates	Early Bird 31-12.2024	Regular 31-1-2025	SPOT REG Feb 2025
Conference Only	6000	7500	9000
Conference + GC Workshop	8000	9500	12000
Conference + 1 Lab Workshop NGS or Functional Genomics	10000	12000	If available

DATES TO REMEMBER
 → EARLY BIRD: 31/12/2024
 → ABSTRACT: 05/02/2025
 Mail 1 page PDF abstract to icgcw.tmc@gmail.com
Best Paper Award for one abstract each in

- Clinical Genetics
- Genetic Counselling
- Molecular Genetics

****Abstracts eligible for fee waiver**

****May Request Reg Fee Refund for abstract with Genotype-Phenotype description of >10 cases of less known genes or VUS**

<< CLICK TO REGISTER >>

Email icgcw.tmc@gmail.com

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PATRON Prof. Sudeep Gupta	CHAIRPERSON Prof. Rajiv Sarin	ORGANIZING SECRETARY Dr Poonam Gera	TREASURER Dr Somya Srivastava
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Less Known or Moderate Risk Genes, Hypomorphic Alleles & VUS

- *Why are they tested in the first place?*
- *Should they be in the report & how should they be described?*
- *Genotype-Phenotype uncertainty of these genes or variants*
- *Family & functional studies to characterise these variants*
- *Pitfalls in ignoring or acting upon these genes / variants*
- ***My RARE GENE DIARY** of Low to Moderate Cancer Risk Genes*

Three Pre Conference Workshops

1) CANCER GENETIC COUNSELLING

- *Genetic counselling overview*
- *Drawing informative complex pedigrees*
- *Psychosocial issues in genetic counselling*
- *Counselling for VUS & Moderate penetrance genes*
- *Counselling for HBOC, Lynch, Li-Fraumeni, NF, FAP, MEN2, Cowden, RB1, Dicer, WT1 & other syndromes*
- *Prenatal & Pre-Implantation genetic testing*
- *Case based discussions & role plays*

Only 40
participants

2) NGS WET LAB, DRY LAB & REPORTING

- *Library prep - Targeted Panel & Exome*
- *Setting up NGS run on MiSeq & NextSeq*
- *Bioinformatics - fastq to vcf, genomic databases*
- *Variant annotation, clinical report drafting on ACMG guidelines. Case based discussions on imp. aspects*
- *Sanger sequencing, Long Range PCR & MLPA*

Only 24
participants

3) FUNCTIONAL GENOMICS & TISSUE CULTURE

- *Functional Genomics Overview*
- *Mammalian Cell Culture Techniques*
- *Gene Editing Technologies - CRISPR/Cas9*
- *Functional Assays in Tissue Culture*
- *DNA Damage Repair and Genomic Integrity*
- *Apoptosis and Cell Survival Assays*

Only 14
participants